



# UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE  
United States Patent and Trademark Office  
Address: COMMISSIONER FOR PATENTS  
P.O. Box 1450  
Alexandria, Virginia 22313-1450  
www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
-----------------	-------------	----------------------	---------------------	------------------

10/590,351

06/20/2007

Gerard Hillion

PET-2271

8163

23599 7590 03/12/2012  
MILLEN, WHITE, ZELANO & BRANIGAN, P.C.  
2200 CLARENDON BLVD.  
SUITE 1400  
ARLINGTON, VA 22201

EXAMINER

PO, MING CHEUNG

ART UNIT

PAPER NUMBER

1771

NOTIFICATION DATE

DELIVERY MODE

03/12/2012

ELECTRONIC

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

docketing@mwzb.com

## Office Action Summary

Application No.

10/590,351

Applicant(s)

HILLION ET AL.

Examiner

MING CHEUNG PO

Art Unit

1771

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

### Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

### Status

- 1) ☒ Responsive to communication(s) filed on 28 September 2011.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ An election was made by the applicant in response to a restriction requirement set forth during the interview on \_\_\_\_; the restriction requirement and election have been incorporated into this action.
- 4) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

### Disposition of Claims

- 5) ☒ Claim(s) 1,9-11,15-19 and 21-29 is/are pending in the application.
- 5a) Of the above claim(s) \_\_\_\_ is/are withdrawn from consideration.
- 6) ☐ Claim(s) \_\_\_\_ is/are allowed.
- 7) ☒ Claim(s) 1,9-11,15-19,23 and 26 is/are rejected.
- 8) ☒ Claim(s) 21-22, 24-25, 27-29 is/are objected to.
- 9) ☐ Claim(s) \_\_\_\_ are subject to restriction and/or election requirement.

### Application Papers

- 10) ☐ The specification is objected to by the Examiner.
- 11) ☐ The drawing(s) filed on \_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 12) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

### Priority under 35 U.S.C. § 119

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
  - ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_.
  - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

### Attachment(s)

- |   |   |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)         | 4) <input type="checkbox"/> Interview Summary (PTO-413)           |
| 2) <input type="checkbox"/> Notice of Draftperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. ____.                                      |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)         | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date ____.   | 6) <input type="checkbox"/> Other: ____.                          |

## DETAILED ACTION

### *Response to Amendment*

1. This is the response to amendment filed for application 10/590351.
2. Claims 1, 9-11, 15-19, 21-29 are currently pending and have been fully considered.

### *Claim Rejections - 35 USC § 103*

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

4. The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
  2. Ascertaining the differences between the prior art and the claims at issue.
  3. Resolving the level of ordinary skill in the pertinent art.
  4. Considering objective evidence present in the application indicating obviousness or nonobviousness.
5. Claims 1, 11, and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over BRADIN (U.S. 5,578,090) in view of GINOSAR et al. (U.S. 6,887, 283).

Regarding claim 1, BRADIN teaches an alternate fuel composition that includes a fuel additive composition. The fuel additive composition is prepared by esterifying the free fatty acids and etherifying glycerol with one or more olefins in the presence of

Art Unit: 1771

an acid catalyst. In lines 22-28 of column 3, BRADIN teaches that the esterification and etherification reactors may be run in separate reactors. BRADIN teaches in lines 34-41 of column 3 that the fatty acid alkyl esters and the glyceryl ethers can be prepared by any means known to those of skill in the art. Means for preparing fatty acid alkyl esters include **transesterifying** triglycerides with **alcohols** in the presence of an **acid or base catalyst**. The alcohol is taught in lines 16-20 of column 4 to be any **C<sub>1-6</sub> straight, branched, or cyclic alcohol, but preferably ethanol**. The glyceryl ethers are prepared by reacting glycerol with an alkyl halide in the presence of a base of an olefin or an alcohol in the presence of an acid catalyst. The olefin is taught in lines 62-67 of column 4 and lines 1-6 of column 5 to be preferably an unsaturated straight, branched, or cyclic hydrocarbon of C<sub>2</sub> to C<sub>10</sub>.

BRADIN does not seem to explicitly teach a heterogeneous catalyst.

However, GINOSAR et al. teaches a process for producing alkyl esters useful in biofuels by transesterifying glyceride or esterifying free fatty acid-containing substances in a single critical phase medium. The transesterification reaction is taught in lines 60-67 of column 4 and lines 1-9 of column 5 that the catalyst used in the transesterification catalyst may be a solid catalyst such as titanium dioxide.

It would be obvious to one of ordinary skill in the art to use the titanium dioxide that GINOSAR et al. teach as the catalyst as the transesterification catalyst in the process that BRADIN teaches.

BRADIN teach in lines 35-43 of column 9 that the catalyst used may be a solid catalyst.

BRADIN does not seem to teach a purification step consisting of a vacuum treatment to remove the ethanol.

However, it would be obvious to one of ordinary skill in the art to use ethanol with the catalyst that GINOSAR et al. teach for the transesterification of fatty acids and an olefin with an acid catalyst for the etherification step.

The motivation to do so can be found in lines 28-31 of column 4 of BRADIN. BRADIN teaches that using olefins rather than alcohol would be less expensive.

A vacuum treatment to an alcohol is well known in the art and it would be obvious for one of ordinary skill in the art to use vacuum treatment to separate the ethanol in the transesterification step from the glycerol before etherification.

Therefore, the invention as a whole would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Regarding claim 11, BRADIN teaches in lines 62-67 of column 4 that **isobutylene (isobutene)** may be used as the olefin in the etherification reaction.

6. Claims 9, 10 and 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over BRADIN (U.S. 5,578,090) in view of GINOSAR et al. (U.S. 6,887, 283) and BOURNAY (U.S. 6,878,837).

The above discussion of BRADIN in view of GINOSAR et al. is incorporated herein by reference. Modified BRADIN does not seem to explicitly teach the conditions of the reactor.

Art Unit: 1771

However, BOURNAY teaches that alkyl esters of fatty acids and high purity glycerin can be produced by using a process comprising a set of transesterification reactions between a vegetable or animal oil and an aliphatic monoalcohol employing a heterogeneous catalyst. BORUNARY teaches the conditions in lines 8-65 of column 4: upflow reactor;  $30 \times 10^5$  to  $80 \times 10^5$  Pa; 453 to 493 K; HSV of  $1.2 \text{ h}^{-1}$  to  $0.1 \text{ h}^{-1}$ . At least 90% by weight of the oil is converted. More than one reactors may be used. The mixture after reaction undergoes a depressurization phase. In lines 1-6 of column 5, the liquid is decanted in a decanter drum.

It would be obvious to one of ordinary skill in the art to apply the conditions that BORUNARY teaches with a reasonable expectation of success given that both BRADIN and BOURNAY are directed towards the production of esters from fatty acids.

The motivation to use the method that BOURNAY teaches can be found in lines 49-59 of column 2 in BOURNAY. BOURNAY teaches that high purity of glycerin can be formed.

Although BOURNAY does not seem to explicitly teach the ranges claimed in the present invention it would be obvious to one of ordinary skill in the art since it has been held that where the general conditions are known, optimization or workable ranges involve only routine experimentation to one of ordinary skill in the art. See *In re Aller*, 220 F.2d 454, 456, 105 USPQ 233, 235 (CCPA 1955).

Therefore, the invention as a whole would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention.

7. Claims 1, 16-19, and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over DELGADO PUCHE (USPGPUB 2003/0167681) and GINOSAR et al. (U.S. 6,887,283).

Regarding claims 1, 16, and 23, DELGADO PUCHE teaches a procedure to produce biodiesel fuels with improved properties at low temperature by **transesterify triglycerides with an alcohol, preferentially methanol or ethanol, in the presence of acid or base catalysts to produce mixtures of methyl or ethyl esters of fatty acids and crude glycerine; isolate the crude glycerin obtained as a secondary product; and then to make all or part of the glycerin react with aldehydes, ketones, to obtain the corresponding acetals.**

DELGADO PUCHE does not seem to explicitly teach a heterogeneous catalyst. However, GINOSAR et al. teaches a process for producing alkyl esters useful in biofuels by transesterifying glyceride or esterifying free fatty acid-containing substances in a single critical phase medium. The transesterification reaction is taught in lines 60-67 of column 4 and lines 1-9 of column 5 that the catalyst used in the transesterification catalyst may be a solid catalyst such as titanium dioxide. It would be obvious to one of ordinary skill in the art to use titanium oxide as the catalyst that GINOSAR et al. as the transesterification catalyst in the process that DELGADO PUCHE teaches.

GINOSAR et al. teach that titanium oxide is a known equivalent for catalysts used for transesterification.

Therefore, the invention as a whole would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Regarding claims 17 and 18, DELGADO PUCHE teaches in paragraph 8 that the glycerine acetals mixed with methyl or ethyl esters of fatty acids in **biodiesel fuels**.

Regarding claim 19, DELGADO PUCHE teaches in paragraph 30 an example reacting **glycerine with acetone**. Glycerine reacts with acetone to form 2,2-dimethyl-1,3,-dioxolane-4-methanol.

8. Claim 15 is rejected under 35 U.S.C. 103(a) as being unpatentable over DELGADO PUCHE (USPGPUB 2003/0167681) and GINOSAR et al. (U.S. 6,887,283) and further in view of NAKAGUCHI (U.S. 3,714,202).

The above discussion of DELGADO PUCHE and GINOSAR et al. is incorporated herein by reference. DELGADO PUCHE does not seem to explicitly state using an acid catalyst in the acetalization step.

However, NAKAGUCHI teaches in lines 22-25 of column 8 that acetal synthesis may be performed with an acid catalyst.

It would be obvious to one of ordinary skill in the art to use an acid catalyst in the acetalization step in the process that DELGADO PUCHE teaches.

The motivation to do so would be to speed up the reaction by use of a catalyst.



Therefore, the invention as a whole would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

***Response to Arguments***

9. Applicant's arguments with respect to claims 1, 9-10, 11, 15-19 have been considered but are moot because the arguments do not apply to any of the references being used in the current rejection.

***Allowable Subject Matter***

10. Claims 22-22, 24-25, 27-29 objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

11. The following is a statement of reasons for the indication of allowable subject matter: The prior art of record do not teach the heterogeneous catalysts that are cited besides titanium oxide.

***Conclusion***

Any inquiry concerning this communication or earlier communications from the examiner should be directed to MING CHEUNG PO whose telephone number is (571)270-5552. The examiner can normally be reached on 9:00 - 4:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Marcheschi can be reached on (571)-272-1374. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Art Unit: 1771

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Ming Cheung Po/  
Patent Examiner  
AU :1771

/Ellen M McAvoy/  
Primary Examiner, Art Unit 1771